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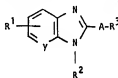
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Fused imidazole compounds, processes for the preparation thereof and pharmaceutical compositions containing them.

The present invention provides new fused imidazole compounds of the formula:



wherein A is a lower alkylene radical, R¹ is a hydrogen or halogen atom or a lower alkyl or lower alkoxy radical, R² is a hydrogen atom or a lower alkyl, cyclo(lower)alkyl, pyridyl, ar(lower)alkyl, which may be substituted by halogen, or anyl radical, which may be substituted by lower alkyl, lower alkoxy, hydroxy or halogen, R³ is an N-containing unsaturated heterocyclic radical, which may be substituted by lower alkyl or amino, and Y is =C- or =N-, and the pharmaceutically-acceptable salts thereof; and processes for the preparation thereof, as well as pharmaceutical compositions comprising them.

These new compounds and the salts thereof are useful as anti-ulcer agents.

FUSED IMIDAZOLE COMPOUNDS, PROCESSES FOR
THE PREPARATION THEREOF AND PHARMACEUTICAL
COMPOSITIONS CONTAINING THEM

The present invention is concerned with new fused
5 imidazole compounds and with the pharmaceutically-
acceptable salts thereof. More particularly, it is
concerned with new fused imidazole compounds and with
the pharmaceutically-acceptable salts thereof which
have an inhibitory activity on ulcers, with processes
10 for the preparation thereof, with pharmaceutical com-
positions containing them and with a method of using
them therapeutically in the treatment of ulcers in
humans and animals.

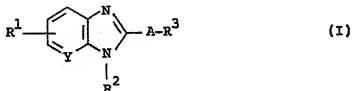
Accordingly, one object of this invention is to
15 provide new fused imidazole compounds and pharmaceutic-
ally-acceptable salts thereof, which are therapeutically
useful for the treatment of ulcers.

Another object of the present invention is to
provide processes for the preparation of these fused
20 imidazole compounds and of the pharmaceutically-
acceptable salts thereof.

A further object of the present invention is to provide pharmaceutical compositions comprising, as an active ingredient, one of the new fused imidazole compounds or a pharmaceutically acceptable salt thereof.

- 5 Yet another object of the present invention is to provide a method of using the new fused imidazole compounds or the pharmaceutically-acceptable salt thereof in the treatment of ulcers in humans and animals.

10 The new fused imidazole compounds of the present invention are compounds of the general formula (I):



wherein A is a lower alkylene radical,

- R^1 is a hydrogen or halogen atom or a lower alkyl or lower alkoxy radical,
- 15 R^2 is a hydrogen atom or a lower alkyl, cyclo-(lower)alkyl, pyridyl, ar(lower)alkyl, which may be substituted by halogen, or aryl radical, which may be substituted by lower alkyl, lower alkoxy, hydroxy or halogen,
- 20 R^3 is an N-containing unsaturated heterocyclic radical, which may be substituted by lower alkyl or amino, and

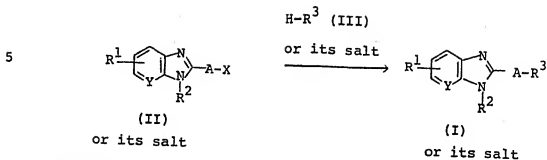
Y is =C- or =N- ;

and the pharmaceutically-acceptable salts thereof.

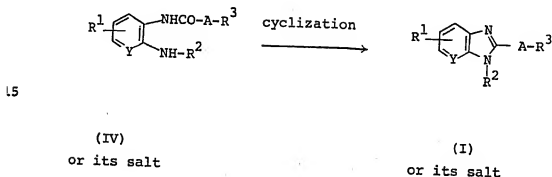
- 25 According to the present invention, the new compounds (I) and the salts thereof can be prepared by the following processes:

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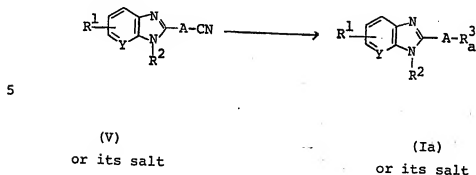
Process 1



Process 2



Process 3



5

wherein R^1 , R^2 , R^3 , A and Y are each as defined above,
 R_a^3 is 1H-tetrazol-5-yl or 2,4-diamino-1,3,5-triazin-6-yl, and X is an acid residue.

In the above and subsequent description of the present specification, suitable examples and illustrations

for the various definitions to be included within the scope of the invention are explained in detail as follows:

The term "lower" is intended to mean 1 to 6 carbon atom(s) unless otherwise indicated.

5 Suitable examples of lower alkylene for A may be methylene, ethylene, ethylidene, propylene, trimethylene, tetramethylene, pentamethylene, hexamethylene or the like.

 Suitable examples of lower alkyl for R¹ and R² may be a straight or branched one such as methyl, ethyl, propyl, 10 isopropyl, butyl, isobutyl, tert-butyl, pentyl, hexyl or the like.

 Suitable examples of lower alkoxy for R¹ may be methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, tert-butoxy, pentyloxy, hexyloxy or the like.

15 Suitable examples of halogen for R¹ may be chlorine, bromine, iodine or fluorine.

 Suitable examples of cyclo(lower)alkyl for R² may be cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl or the like.

20 Suitable examples of ar(lower)alkyl for R² may be benzyl, phenethyl, 3-phenylpropyl, benzhydryl, trityl or the like. Said ar(lower)alkyl group may be substituted with the aforementioned halogen, and suitable examples of the halogenated ar(lower)alkyl may be 2-chlorobenzyl, 25 4-chlorobenzyl, 4-bromobenzyl, 4-iodobenzyl, 4-fluorobenzyl, 2,4-dichlorobenzyl, 2-bromo-4-chlorobenzyl, 4-chlorophenethyl, 3-(4-chlorophenyl)propyl, 4-chloro-benzhydryl, 4,4'-dichlorobenzhydryl or the like.

 Suitable examples of aryl for R² may be phenyl, 30 naphthyl or the like. Said aryl group may have one or more substituent(s) selected from the groups consisting of hydroxy and the aforementioned lower alkyl, lower alkoxy and halogen.

 Suitable examples of aryl group having such 35 substituent(s) may be lower alkyl substituted aryl [e.g.

tolyl, xylyl, 4-ethylphenyl, 4-methyl-1-naphthyl, etc.], lower alkoxy substituted aryl [e.g. 4-methoxyphenyl, 3,4-dimethoxyphenyl, 6-ethoxyphenyl, 5-methoxy-1-naphthyl, etc.], halogenated aryl [e.g. 3-chlorophenyl, 4-chlorophenyl, 2,4-dichlorophenyl, 8-chloro-2-naphthyl, etc.], hydroxy substituted aryl [e.g. 2-hydroxyphenyl, 4-hydroxyphenyl, 1-hydroxy-2-naphthyl, etc.] or the like.

Suitable examples of N-containing unsaturated heterocyclic group for R³ may be unsaturated 3- to 9-membered (preferably 5- to 6-membered) mono or polycyclic heterocyclic group containing 1 to 4 nitrogen atom(s) such as imidazolyl, pyrazolyl, triazolyl [e.g. 1H-1,2,3-triazolyl, 1H-1,2,4-triazolyl, 2H-1,2,3-triazolyl, 4H-1,2,4-triazolyl, etc.], tetrazolyl [e.g. 1H-tetrazolyl, 2H-tetrazolyl, etc.], pyridyl [e.g. 2-pyridyl, 3-pyridyl or 4-pyridyl], pyrimidinyl, pyrazinyl, pyridazinyl, 1,3,5-triazinyl, 2-oxo-benzothiazolinyl, or the like.

The above-mentioned heterocyclic groups may be substituted with the aforementioned lower alkyl group(s) or amino group(s). Suitable examples of the heterocyclic group having such substituent(s) may be imidazolyl [e.g. 2-methylimidazol-1-yl, 4-methylimidazol-1-yl, 5-methylimidazol-1-yl, 4-ethylimidazol-1-yl, 4-hexylimidazol-1-yl, etc.], pyrazolyl [e.g. 3-methylpyrazol-1-yl, 4-methylpyrazol-1-yl, etc.], triazolyl [e.g. 4-methyl-1H-1,2,3-triazol-1-yl, 3-methyl-1H-1,2,4-triazol-1-yl, 4-methyl-2H-1,2,3-triazol-2-yl, 3-methyl-4H-1,2,4-triazol-4-yl, etc.], tetrazolyl [e.g. 5-methyl-1H-tetrazol-1-yl, 1-methyl-1H-tetrazol-5-yl, 2-methyl-2H-tetrazol-5-yl, etc.], pyridyl [e.g. 4-methylpyridin-3-yl, etc.], pyrimidinyl [e.g. 5-methylpyrimidin-2-yl, etc.], pyrazinyl [e.g. 5-methylpyrazin-2-yl, etc.], pyridazinyl [e.g. 6-methylpyridazin-3-yl, etc.], 2,4-diamino-1,3,5-triazin-6-yl, 5-methyl-2-oxo-benzothiazolin-3-yl, or the like.

Suitable examples of the acid residue for X may be halide [e.g. chloride, bromide, iodide, etc.], sulfonate [e.g. benzenesulfonate, tosylate, etc.] or the like.

5 Suitable pharmaceutically acceptable salts of the object compounds (I) are conventional non-toxic salts and include an organic acid salt [e.g. formate, acetate, trifluoroacetate, maleate, fumarate, tartrate, methanesulfonate, benzenesulfonate, toluenesulfonate, etc.],
10 an inorganic acid salt [e.g. hydrochloride, hydrobromide, sulfate, phosphate, etc.], a salt with an amino acid [e.g. arginine salt, glutamic acid salt, ornithine salt, etc.] or the like.

15 The processes for preparing the object compounds (I) and salts thereof are explained in detail in the following.
Process 1

 The object compound (I) and its salt can be prepared by reacting a compound (II) or its salt with a compound
20 (III) or its salt.

 Suitable salts of the compounds (II) and (III) may be the same as those exemplified for the compound (I).

 This reaction is usually carried out in the presence of a base.

25 Suitable base may include an inorganic base such as alkali metal hydride [e.g. sodium hydride, potassium hydride, etc.], alkali metal hydroxide [e.g. sodium hydroxide, potassium hydroxide, etc.], alkaline earth metal hydroxide [e.g. magnesium hydroxide, calcium hydroxide, etc.],
30 alkali metal carbonate [e.g. sodium carbonate, potassium carbonate, etc.], alkaline earth metal carbonate [e.g. magnesium carbonate, calcium carbonate, etc.], alkali metal bicarbonate [e.g. sodium bicarbonate, potassium bicarbonate, etc.], alkali metal acetate [e.g. sodium
35 acetate, potassium acetate, etc.], alkaline earth metal

phosphate [e.g. magnesium phosphate, calcium phosphate, etc.], alkali metal hydrogen phosphate [e.g. disodium hydrogen phosphate, dipotassium hydrogen phosphate, etc.], and an organic base such as trialkylamine [e.g. trimethylamine, triethylamine, etc.], picoline, N-methylpyrrolidine, N-methylmorpholine. And further, compound (III) per se can also be used as a base.

This reaction is usually carried out in a solvent such as alcohol [e.g. methanol, ethanol, etc.], benzene, N,N-dimethylformamide, tetrahydrofuran, diethyl ether or any other solvent which does not adversely affect the reaction.

The reaction temperature is not critical and the reaction is usually carried out at ambient temperature, or under warming or heating.

In case that the compound (II) possesses two or more reactive sites in the molecule, this reaction is preferably carried out by the following steps in order to increase the selectivity of the reactive site.

- 1) reacting a compound (II) or its salt with a compound (III) protected by a conventional protective group [e.g. formyl, acetyl, etc.] at one or more reactive site(s) or its salt,
- 2) eliminating the protective group(s) of the reaction product by a conventional manner [e.g. hydrolysis, etc.].

This method can be carried out according to Example mentioned later or analogous method thereto, and this method is also included within the scope of Process 1.

Process 2

The object compound (I) and its salt can be prepared by cyclizing the compound (IV) or its salt.

Suitable salts of the compound (IV) may be the same as those exemplified for the compound (I).

This reaction may be preferably carried out in the presence of an acid such as inorganic acid [e.g. hydrochloric acid, sulfuric acid, polyphosphoric acid, etc.], organic acid [e.g. trifluoroacetic acid, benzenesulfonic acid, toluenesulfonic acid, etc.] or the like.

This reaction can be carried out in the absence or presence of a conventional solvent such as aromatic hydrocarbon [e.g. benzene, toluene, xylene, etc.], alcohol [e.g. methanol, ethanol, propanol, isopropyl alcohol, butanol, tert-butyl alcohol, etc.] or any other organic solvent which does not adversely influence the reaction. In case that the acid to be used is liquid, it can also be used as a solvent.

This reaction is preferably carried out under dehydrating condition such as an azeotropic dehydration, in the presence of a dehydrating agent [e.g. anhydrous magnesium sulfate, anhydrous zinc chloride, phosphorus pentoxide, zeolite, silica gel, etc.], or the like.

The reaction temperature is not critical, and the reaction is usually carried out under warming or heating.

Process 3

The object compound (Ia) and its salt can be prepared by reacting a compound (V) or its salt with dicyanodiamide or its salt or an azide compound.

Suitable salts of dicyanodiamide may be the same as those exemplified for the compound (I).

Suitable example of the azide compound may be an inorganic base salt of azide [e.g. sodium azide, potassium azide, lithium azide, calcium azide, barium azide, etc.], hydrogen azide, hydrazonic acid, ammonium azide or the like.

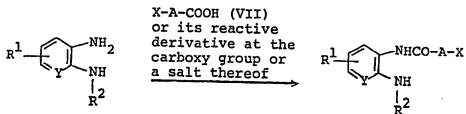
This reaction is usually carried out in a conventional solvent which does not adversely influence the reaction

such as water, methanol, ethanol, dioxane, N,N-dimethylformamide, methyl cellosolve, ethyl cellosolve or a mixture thereof.

The reaction temperature is not critical, and the reaction is usually carried out at ambient temperature or under warming or heating.

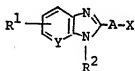
Among the starting compounds (II) and (IV), the new compounds can be prepared by the following processes.

Process A



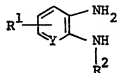
(VI)
or its reactive derivative
at the amino group or a
salt thereof

(VIII)
or its salt
cyclization



(II)
or its salt

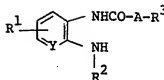
Process B



(VI)

or its reactive derivative
at the amino group or a
salt thereof

$\text{R}^3\text{-A-COOH}$ (IX)
or its reactive
derivative at the
carboxy group or
a salt thereof



(IV)

or its salt

In the above formulas, R^1 , R^2 , R^3 , A, Y and X are
each as defined above.

The processes for preparing the starting compounds
(II) and (IV) and salts thereof are explained in detail
in the following.

Process A

i) Preparation of the compound (VIII) (Step 1):

The compound (VIII) and its salt can be prepared by
reacting a compound (VI) or its reactive derivative at
the amino group or a salt thereof with a compound (VII)
or its reactive derivative at the carboxy group or a
salt thereof.

Suitable salts of the compound (VI) may be the same
as those exemplified for the compound (I), and suitable
salts of the compound (VII) may be a conventional organic
or inorganic base addition salt [e.g. sodium salt,
potassium salt, triethylamine salt, etc.].

Suitable reactive derivatives at the amino group of
the compound (VI) include conventional ones used in
amidation, for example, Schiff's base type imino or its
tautomeric enamine type isomer formed by reaction of

the compound (VI) with a carbonyl compound, a silyl derivative formed by reaction of the compound (VI) with a silyl compound such as trimethylsilylacetamide, bis(trimethylsilyl)acetamide or the like, a derivative
5 formed by reaction of the compound (VI) with phosphorus trichloride or phosgene, and the like.

Suitable reactive derivative at the carboxy group of the compound (VII) may include an acid halide, an acid anhydride, an ester, an activated amide, an activated
10 ester and the like.

Suitable examples of such reactive derivatives may be an ester such as lower alkyl ester [e.g. methyl ester, ethyl ester, propyl ester, hexyl ester, etc.], acid chloride, an acid azide, a mixed acid anhydride with
15 an acid such as substituted phosphoric acid [e.g. dialkylphosphoric acid, phenylphosphoric acid, etc.], aliphatic carboxylic acid [e.g. pivalic acid, acetic acid, trichloroacetic acid, etc.] or the like, a symmetrical acid anhydride, an activated amide with
20 imidazole, triazole or dimethylpyrazole, an activated ester with N-hydroxysuccinimide, N-hydroxyphthalimide or 1-hydroxy-6-chlorobenzotriazole, and the like.

The reactive derivatives of the compounds (VI) and (VII) can be selected according to the kinds of the
25 compounds (VI) and (VII), respectively.

When the compound (VII) is used in a free acid form or its salt form in the reaction, the reaction is preferably carried out in the presence of a conventional condensing agent such as N,N'-dicyclohexylcarbodiimide,
30 N-cyclohexyl-N'-morpholinoethylcarbodiimide, N-ethyl-N'-(3-dimethylaminopropyl)carbodiimide, thionyl chloride, phosphorus oxychloride, phosphorus oxybromide, oxalyl chloride, lower alkoxy carbonyl halide [e.g. ethyl chloroformate, isobutyl chloroformate, etc.], 1-(p-chlorobenzenesulfonyloxy)-6-chloro-1H-benzotriazole, or the
35 like.

The reaction is usually carried out in a conventional solvent such as water, acetone, dioxane, chloroform, methylene chloride, ethylene chloride, tetrahydrofuran, ethyl acetate, N,N-dimethylformamide, pyridine or any other organic solvent which does not adversely influence the reaction. Among these solvents, hydrophilic solvents may be used in a mixture with water.

The reaction may be preferably carried out in the presence of an inorganic or an organic base such as an alkali metal hydroxide [e.g. sodium hydroxide, potassium hydroxide, etc.], an alkali metal carbonate [e.g. sodium carbonate, potassium carbonate, etc.], an alkali metal bicarbonate [e.g. sodium bicarbonate, potassium bicarbonate, etc.], tri(lower)alkylamine [e.g. trimethylamine, triethylamine, etc.], pyridine or its derivative [e.g. picoline, lutidine, 4-dimethylamino-pyridine, etc.], or the like. In case that the base or the condensing agent to be used is liquid, it can also be used as a solvent.

The reaction temperature is not critical, and the reaction can be carried out under cooling, at ambient temperature or under warming or heating.

ii) Preparation of the compound (II) (Step 2):

The compound (II) and its salt can be prepared by cyclizing a compound (VIII) or its salt.

This reaction can be carried out substantially in the same manner as Process 2, and therefore the reaction mode and reaction conditions [e.g. acid, dehydrating agent, solvent, reaction temperature, etc.] of this reaction are to be referred to those as explained in Process 2.

Process B

The compound (IV) and its salt can be prepared by

reacting a compound (VI) or its reactive derivative at the amino group or a salt thereof with a compound (IX) or its reactive derivative at the carboxy group or a salt thereof.

5 Suitable salts of the compound (VI) may be the same as those exemplified for the compound (I), and suitable salts of the compound (IX) may be a conventional organic or inorganic base addition salt [e.g. sodium salt, potassium salt, triethylamine salt, etc.].

10 This reaction can be carried out substantially in the same manner as that of Step 1 of Process A, and therefore the reaction mode and reaction conditions [e.g. reactive derivatives, condensing agent, solvent, reaction temperature, etc.] of this reaction are to be referred to
15 those as explained in Step 1 of Process A.

The reaction product can be used in the next step (i.e. Process 2) with or without isolation. The isolation can be carried out by a conventional manner.

20 It is to be noted that each of the object compound (I) and the starting compounds (II), (IV) and (V) include one or more stereoisomers due to asymmetric carbon atoms in the molecule, and all of such isomers of the compounds (I), (II), (IV) and (V) are included within the scope of
25 this invention.

The object compounds (I) and their pharmaceutically acceptable salts of the present invention are novel and exhibit high inhibitory activity on ulcer.

30 In order to illustrate the usefulness of the object compound (I), the pharmacological data of some of the object compounds (I) are shown in the following.

(A) Inhibition on ethanol ulcer

35 Test Method :

Five male Sprague-Dawley rats, aged 7 weeks and weighing about 200 g, were used per group for the study on ethanol ulcer after the fast for 24 hours.

5 Test compound was suspended in 0.1% methylcellulose aqueous solution, and the suspension (5 ml/kg) was orally given to each rat.

The control group was given a vehicle, i.e. 0.1% methylcellulose aqueous solution (5 ml/kg), alone in the same way.

10 Absolute ethanol (5 ml/kg) was orally administered 30 minutes after dosing with test compound, and one hour later, the rats were sacrificed and their stomachs were removed. The area of ulcers of each rat was measured. The mean area (mm^2) in the medicated group was compared
15 with that in the control group.

Test Result:

Test Compound: 2-(Imidazol-1-yl)methyl-1-phenyl-1H-benzimidazole

20 The ED_{50} value of the test compound: 3.4 mg/kg

(B) Inhibition on stress ulcer

Test Method:

Five Sprague-Dawley rats weighing about 200 g were used per group. Each animal was immobilized in a small cage and put in a water bath allowing to respire. The temperature of the water bath kept at 22°C. The test compound was administered orally just before the immobilization. Seven hours later, the animals were
30 sacrificed and their stomachs were removed. The stomach was then fixed with 2% formalin. The area of ulcers was measured for each animal. The mean area (mm^2) in the medicated animals was compared with that in the control animals.
35

Test Result:

Test Compound: 2-(Imidazol-1-yl)methyl-1-phenyl-1H-benzimidazole

The ED₅₀ value of the test compound: 7.9 mg/kg

(C) Gastric secretion in Heidenhain pouch dogs

Test Method:

Beagle dogs, weighing about 8-13 kg, were used for the study on gastric secretion. The animals were surgically provided with a vagally denervated Heidenhain pouch. One month or more later, the dogs were fasted overnight. Gastric secretion was stimulated by an intravenous infusion of tetragastrin (10 µg/kg/hr). Gastric samples were collected at 15 min intervals. After its volume was almost constant, test compound suspended in 0.1% methylcellulose solution was injected intravenously (0.2 ml/kg). Acid concentration was determined by titrating an aliquot to pH 7.0 with 0.1N sodium hydroxide solution using automatic titration (Hiranuma RAT-11 Type). Total acid output was calculated by multiplying total volume of gastric samples by acid concentration, and percentage change of total acid output was calculated by comparing with predosing value of test compound.

Test Result:

Test Compound: 2-(Pyridin-3-yl)methyl-1-phenyl-1H-benzimidazole

The inhibition of the test compound (Dose: 1 mg/kg, iv): 92.9%

As being apparent from the above test results, the object compounds [I] of the present invention are useful as antiulcer medicines.

For therapeutic purpose, the compounds according to the present invention can be used in a form of pharmaceutical preparation containing said compound as an active ingredient, in admixture with a pharmaceutically acceptable carrier such as an organic or inorganic solid or liquid excipient suitable for oral or parenteral administration. The pharmaceutical preparations may be capsules, tablets, dragees, solution, suspension, emulsion, and the like. If desired, there may be included in the above preparations auxiliary substances, stabilizing agents, wetting or emulsifying agents, buffers and other commonly used additives.

While the dosage of the compounds will vary depend upon the age and condition of the patient, an average single dose of about 5 mg, 10 mg, 50 mg, 100 mg, 250 mg, 500 mg, and 1000 mg of the compounds according to the present invention may be effective for treating ulcer. In general, amounts between 1 mg/body and about 6,000 mg/body or even more may be administered per day.

The following preparations and examples are given for the purpose of illustrating the present invention.

Preparation 1

To a solution of 2-anilinoaniline (4.0 g) in a mixture of dichloromethane (20 ml) and pyridine (6.87 g) was dropwise added 4-chlorobutyl chloride (4.29 g) at 8°C during a period of 16 minutes. The mixture was stirred at the same temperature for 40 minutes and then diluted with ice-water. The mixture was extracted twice with dichloromethane. The combined extract was washed with 1N hydrochloric acid and water in turn and dried over magnesium sulfate. The solvent was removed under reduced pressure to give a crystalline residue of 4-chloro-N-(2-anilinophenyl)butyramide (4.3 g).

IR (Nujol) : 3350, 3200, 1635, 1585 cm^{-1}

NMR (CDCl_3 , δ) : 1.8-2.65 (4H, m), 3.57 (2H, t, $J=6\text{Hz}$), 5.63 (1H, m), 6.67-8.17 (10H, m)

Preparation 2

2-Chloro-N-(2-anilinophenyl)propanamide was obtained according to substantially the same manner as that of Preparation 1.

mp 73-78°C

IR (Film) : 3350, 1670, 1610 cm^{-1}

NMR (CDCl_3 , δ) : 1.7 (3H, d, $J=7\text{Hz}$), 4.45 (1H, q, $J=7\text{Hz}$), 4.8 (1H, m), 6.65-7.5 (8H, m), 7.8-8.15 (1H, m), 8.7 (1H, m)

Preparation 3

2-Chloro-N-[2-(3-methylanilino)phenyl]acetamide was obtained according to substantially the same manner as that of Preparation 1.

mp 112-113°C

IR (Nujol) : 3350, 3315, 1670, 1595 cm^{-1}

NMR (CDCl_3 , δ) : 2.27 (3H, s), 4.1 (2H, s), 5.4 (1H, m), 6.37-7.4 (7H, m), 7.8-8.2 (1H, m), 8.65 (1H, m)

Preparation 4

2-Chloro-N-[2-(4-methoxyanilino)phenyl]acetamide
was obtained according to substantially the same manner
as that of Preparation 1.

5

mp 93-94°C

IR (Nujol) : 3325, 3220, 1640, 1600 cm^{-1}

NMR (CDCl_3 , δ) : 3.75 (3H, s), 4.12 (2H, s),
5.4 (1H, m), 6.8-7.3 (7H, m), 7.7-8.05 (1H, m),
8.65 (1H, m)

10

Preparation 5

2-Chloro-N-[2-(4-chloroanilino)phenyl]acetamide
was obtained according to substantially the same manner
as that of Preparation 1.

15

mp 96-97°C

IR (Nujol) : 3360, 3250, 1660, 1605 cm^{-1}

NMR (CCl_4 , δ) : 4.02 (2H, s), 5.65 (1H, m),
6.55-7.28 (7H, m), 7.65-8.02 (1H, m),
8.55 (1H, m)

20

Preparation 6

2-Chloro-N-(5-methoxy-2-anilinophenyl)acetamide
was obtained according to substantially the same manner
as that of Preparation 1.

25

mp 94-95°C

IR (Nujol) : 3380, 3340, 1665, 1585 cm^{-1}

NMR (CDCl_3 , δ) : 3.8 (3H, s), 4.03 (2H, s),
5.1 (1H, m), 6.45-7.4 (7H, m), 7.93 (1H, d,
J=3Hz), 8.97 (1H, m)

30

Preparation 7

A solution of 2-chloro-N-(2-anilinophenyl)acetamide
(10 g) and p-toluenesulfonic acid monohydrate (4.5 g)
in benzene (250 ml) was refluxed for 1.5 hours under an
azeotropic dehydration. After being cooled to ambient

35

temperature, the reaction mixture was diluted with an aqueous sodium bicarbonate solution and extracted with ethyl acetate three times. The combined organic layer was washed successively with water and brine, dried over anhydrous magnesium sulfate and evaporated under reduced pressure. The residual oil was purified by column chromatography on silica gel (100 g) with dichloromethane as an eluent to afford 2-chloromethyl-1-phenyl-1H-benzimidazole (3.4 g).

mp 117-119°C

IR (Nujol) : 1610, 1592, 1495 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 4.64 (2H, s), 7.0-7.9 (9H, m)

Preparation 8

2-(1-Chloroethyl)-1-phenyl-1H-benzimidazole was obtained according to substantially the same manner as that of Preparation 7.

mp 71-72°C

IR (Nujol) : 1600, 1585, 1490 cm^{-1}

NMR (CCl_4 , δ) : 2.03 (3H, d, $J=7\text{Hz}$), 4.95 (1H, q, $J=7\text{Hz}$), 6.85-7.9 (9H, m)

Preparation 9

2-Chloromethyl-1-(3-methylphenyl)-1H-benzimidazole was obtained according to substantially the same manner as that of Preparation 7.

IR (Film) : 1600, 1585, 1510 cm^{-1}

NMR (CDCl_3 , δ) : 2.47 (3H, s), 4.7 (2H, s), 7.07-8.07 (8H, m)

Preparation 10

2-Chloromethyl-1-(4-methoxyphenyl)-1H-benzimidazole was obtained according to substantially the same manner as that of Preparation 7.

mp 131-133°C

IR (Nujol) : 1610, 1585, 1515, 1460 cm^{-1}
NMR (CDCl_3 , δ) : 3.9 (3H, s), 4.63 (2H, s),
6.95-8.0 (8H, m)

5 Preparation 11

2-Chloromethyl-1-(4-chlorophenyl)-1H-benzimidazole
was obtained according to substantially the same manner
as that of Preparation 7.

mp 126-127°C

10 IR (Nujol) : 1615, 1590, 1495 cm^{-1}
NMR (CDCl_3 , δ) : 4.67 (2H, s), 6.9-7.95 (8H, m)

Preparation 12

15 2-Chloromethyl-5-methoxy-1-phenyl-1H-benzimidazole
was obtained according to substantially the same manner
as that of Preparation 7.

mp 121-123°C

IR (Nujol) : 1620, 1580, 1495 cm^{-1}
NMR (CDCl_3 , δ) : 3.85 (3H, s), 4.65 (2H, s),
20 6.7-7.73 (8H, m)

Preparation 13

A solution of 4-chloro-N-(2-anilinophenyl)butyramide
(0.5 g) and p-toluenesulfonic acid monohydrate (33 mg) in
25 toluene (15 ml) was refluxed for 1.5 hours. Toluene was
removed under reduced pressure to afford a residue, which
was dissolved in dichloromethane (20 ml), washed with a
saturated aqueous solution of sodium bicarbonate and
water in turn and dried over magnesium sulfate. Removal
30 of the solvent under reduced pressure afforded an oily
residue, which was subjected to column chromatography on
silica gel (5 g) eluting with a mixture of dichloromethane
and ethyl acetate (10:1) to give crystals of 2-(3-
chloropropyl)-1-phenyl-1H-benzimidazole (148 mg).

35 mp 71-72°C

IR (Nujol) : 1610, 1595, 1510, 1500 cm^{-1}

NMR (CDCl_3 , δ) : 1.85-2.7 (2H, m), 2.95 (2H, t,

$J=6\text{Hz}$), 3.5-4.0 (2H, m), 7.0-7.9 (9H, m)

Mass: 270 (M^+)

5

The second fraction was combined and evaporated to give crystals of 2-(3-hydroxypropyl)-1-phenyl-1H-benzimidazole (128 mg).

mp 94-95°C

0

IR (Nujol) : 3300, 1610, 1595, 1500 cm^{-1}

NMR (CDCl_3 , δ) : 1.85-2.7 (2H, m), 2.8-3.2 (2H, m),
3.5-3.9 (2H, m), 7.0-8.0 (9H, m)

Mass : 252 (M^+)

5

Preparation 14

A solution of 2-(3-hydroxypropyl)-1-phenyl-1H-benzimidazole (1.15 g) and thionyl chloride (0.6 g) in chloroform (23 ml) was refluxed for 1.5 hours. After being cooled, the mixture was diluted with a saturated aqueous solution of sodium bicarbonate. The chloroform layer was washed with water and a saturated aqueous solution of sodium chloride in turn, and dried over magnesium sulfate. The solvent was evaporated under reduced pressure to afford 2-(3-chloropropyl)-1-phenyl-1H-benzimidazole (1.0 g).

0

5

IR (Nujol) : 1610, 1595, 1510, 1500 cm^{-1}

Preparation 15

To a mixture of 3-amino-2-anilinopyridine (0.5 g), 2-(imidazol-1-yl)acetic acid (0.38 g) and pyridine (0.24 ml) in chloroform (5 ml) was dropwise added phosphorus oxychloride (0.28 ml) and the mixture was refluxed for 1.5 hours. After being cooled, the reaction mixture was decanted and the residue was poured into saturated aqueous sodium bicarbonate. The mixture was

0

5

extracted three times with ethyl acetate. The combined
extract was dried over magnesiumsulfate and the solvent
was evaporated under reduced pressure. The resultant
residue was triturated in ethyl acetate (10 ml) to give
5 2-(imidazol-1-yl)-N-(2-anilinopyridin-3-yl)acetamide
(200 mg).

mp 151-154°C

IR (Nujol) : 3350, 1700, 1640, 1610 cm^{-1}

NMR (DMSO-d_6 , δ) : 4.95 (2H, s), 6.7-8.3 (12H, m),

9.73 (1H, brs)

Mass : 293 (M^+)

Example 1

A solution of 2-chloromethyl-1-phenyl-1H-benzimidazole (3.4 g) and imidazole (4.8 g) in ethanol (25 ml) was stirred at 60°C for 7 hours. The reaction mixture was concentrated under reduced pressure to give an oil, which was diluted with water and extracted three times with dichloromethane. The combined extract was washed twice with water and once with brine, dried over anhydrous magnesium sulfate and evaporated under reduced pressure. The residue was purified by column chromatography on silica gel (100 g) with a mixture of methanol and dichloromethane (3:97) as an eluent to afford an oil, which was crystallized from a mixture of ethyl acetate and hexane to give 2-(imidazol-1-yl)methyl-1-phenyl-1H-benzimidazole (2.65 g).

mp. 116-117°C

IR (Nujol) : 1612, 1590, 1495 cm^{-1}

NMR ($\text{DMSO}-d_6$, δ) : 5.40 (2H, s), 6.8-7.8 (12H, m)

Analysis: Calcd. for $\text{C}_{17}\text{H}_{14}\text{N}_4$

Calcd.: C 74.43 H 5.14 N 20.43

Found : C 73.84 H 5.14 N 20.25

Example 2

A solution of 2-chloromethyl-1H-benzimidazole (4.85 g) and imidazole (9.9 g) in ethanol (37 ml) was stirred at 70°C for 6 hours. The reaction mixture was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (120 g) with a mixture of methanol and dichloromethane (5:95) as an eluent to afford an oil, which was crystallized from ethyl acetate. Recrystallization from a mixture of methanol and diethyl ether to give 2-(imidazol-1-yl)methyl-1H-benzimidazole (0.76 g).

mp. 198-200°C

IR (Nujol) : 3100, 1660, 1620, 1515 cm^{-1}

NMR (DMSO- d_6 , δ) : 5.14 (2H, s), 6.86 (1H, s),
7.0-7.64 (6H, m), 7.76 (1H, s)

Analysis: Calcd. for $C_{11}H_{10}N_4$

Calcd.: C 66.65 H 5.09 N 28.51

Found : C 66.66 H 5.05 N 28.51

Example 3

A solution of 2-chloromethyl-1-(4-chlorobenzyl)-1H-benzimidazole (2.0 g) and imidazole (2.34 g) in acetonitrile (20 ml) was heated at 90°C for 1.5 hours. The reaction mixture was concentrated under reduced pressure to give an oil, which was diluted with water and extracted three times with ethyl acetate. The combined extract was washed twice with water, dried over anhydrous magnesium sulfate and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (30 g) with a mixture of methanol and dichloromethane (10:90) as an eluent to give an oil (1.6 g). To a solution of the oil in methanol (20 ml) was added fumaric acid (0.59 g). The mixture was filtered to remove an insoluble material and the filtrate was evaporated under reduced pressure to give a crystalline residue, which was recrystallized from methanol to afford pure 2-(imidazol-1-yl)methyl-1-(4-chlorobenzyl)-1H-benzimidazole fumarate (1.6 g).

mp. 162-163°C

IR (Nujol) : 1690, 1635, 1600, 1515, 1495 cm^{-1}

NMR (DMSO- d_6 , δ) : 5.67 (4H, s), 6.67 (2H, s),

7.0-7.8 (10H, m), 7.93 (1H, brs), 9.53 (2H, brs)

Analysis: Calcd. for $C_{22}H_{19}Cl_2N_5O_4$

Calcd.: C 60.21 H 4.36 N 12.77

Found : C 59.65 H 4.37 N 12.78

Example 4

To a solution of imidazole (226 mg) in N,N-

dimethylformamide (9 ml) was added sodium hydride (60%
in oil, 140 mg) with stirring and ice-cooling. The
mixture was stirred for 15 minutes at the same temperature
and for 1 hour at 100°C. To the resultant mixture, after
cooling, was added 2-(3-chloropropyl)-1-phenyl-1H-
benzimidazole (0.9 g) and the mixture was stirred for 1
hour at 100°C. After being cooled, the reaction mixture
was diluted with water and extracted with dichloromethane.
The combined extract was washed with water and saturated
aqueous sodium chloride in turn, and dried over magnesium
sulfate. The solvent was removed under reduced pressure
to give an oily residue, which was subjected to column
chromatography on silica gel (9 g) eluting with a mixture
of dichloromethane and methanol (50:1) to give an oil.
To a solution of the oil in diethyl ether (30 ml) was
added a solution of oxalic acid dihydrate (321 mg) in
ethanol to give an oily precipitate. The oily precipitate
was triturated in ethanol to give powder of 2-[3-(imidazol-1-
yl)propyl]-1-phenyl-1H-benzimidazole oxalate (920 mg).

mp. 141-143°C

IR (Nujol) : 1610, 1590, 1500, 1210 cm^{-1}

NMR (DMSO- d_6 , δ) : 2.0-3.0 (4H, m), 4.25 (2H, t,
J=6.5Hz), 7.0-7.9 (11H, m), 8.6 (1H, s)

Analysis: Calcd. for $\text{C}_{19}\text{H}_{18}\text{N}_4 \cdot \text{C}_2\text{H}_2\text{O}_4$

Calcd.: C 64.28 H 5.14 N 14.28

Found : C 64.37 H 5.19 N 14.31

Example 5

To a solution of sodium ethoxide, prepared from
sodium metal (230 mg) and ethanol (24 ml), were added
1H-1,2,4-triazole (690 mg) and 2-chloromethyl-1-phenyl-
1H-benzimidazole (2.4 g) with stirring at ambient tempera-
ture. The mixture was refluxed for 1.5 hours and
evaporated under reduced pressure. The resultant residue
was subjected to column chromatography on silica gel

(50 g) eluting with a mixture of chloroform and methanol (200:3) to give a crystalline product. The crystal was recrystallized from isopropyl alcohol to give 2-(1H-1,2,4-triazol-1-yl)methyl-1-phenyl-1H-benzimidazole (1.65 g).

mp. 154-156°C

IR (Nujol) : 1595, 1500, 1450 cm^{-1}

NMR (CDCl_3 , δ) : 5.5 (2H, s), 7.0-8.0 (9H, m),
7.83 (1H, s), 8.05 (1H, s)

10 Analysis: Calcd. for $\text{C}_{16}\text{H}_{13}\text{N}_5$

Calcd.: C 69.80 H 4.76 N 25.44

Found : C 69.77 H 4.93 N 25.22

Example 6

15 2-(4-Methylimidazol-1-yl)methyl-1-phenyl-1H-benzimidazole·mono oxalate· $\frac{1}{2}$ hydrate was obtained according to substantially the same manner as that of Example 1.

mp. 137-138°C

IR (Nujol) : 1720, 1630, 1600, 1500 cm^{-1}

20 NMR ($\text{DMSO}-d_6$, δ) : 2.2 (3H, s), 5.62 (2H, s),
7.0-7.9 (10H, m), 8.3 (1H, s), 9.25 (2H, s)

Analysis: Calcd. for $\text{C}_{18}\text{H}_{16}\text{N}_4 \cdot \text{C}_2\text{H}_2\text{O}_4 \cdot \frac{1}{2}\text{H}_2\text{O}$

Calcd.: C 62.00 H 4.94 N 14.46

Found : C 62.53 H 5.05 N 14.25

Example 7

2-(2-Methylimidazol-1-yl)methyl-1-phenyl-1H-benzimidazole was obtained according to substantially the same manner as that of Example 1.

30 mp. 124-126°C

IR (Nujol) : 1610, 1595, 1500 cm^{-1}

NMR (CDCl_3 , δ) : 2.0 (3H, s), 5.15 (2H, s), 6.6
(1H, d, J=2Hz), 6.8 (1H, d, J=2Hz),
6.9-8.0 (9H, m)

35 Analysis: Calcd. for $\text{C}_{18}\text{H}_{16}\text{N}_4$

Calcd. : C 74.98 H 5.59 N 19.43

Found : C 75.07 H 5.71 N 19.47

Example 8

2-[1-(Imidazol-1-yl)ethyl]-1-phenyl-1H-benzimidazole dihydrochloride· $\frac{1}{2}$ hydrate was obtained according to substantially the same manner as that of Example 1.

mp. 175-190°C

IR (Nujol) : 3350, 2550, 2495, 1580, 1545 cm^{-1}

NMR (DMSO- d_6 , δ) : 1.9 (3H, d, J=7Hz), 6.18 (1H, q, J=7Hz), 6.95-8.0 (11H, m), 9.17 (1H, m)

Analysis: Calcd. for $\text{C}_{18}\text{H}_{16}\text{N}_4 \cdot 2\text{HCl} \cdot \frac{1}{2}\text{H}_2\text{O}$

Calcd.: C 58.39 H 5.17 N 15.13

Found : C 58.12 H 5.42 N 14.88

Example 9

2-(Imidazol-1-yl)methyl-1-(3-methylphenyl)-1H-benzimidazole was obtained according to substantially the same manner as that of Example 1.

mp. 91-93°C

IR (Nujol) : 1603, 1590, 1518, 1502 cm^{-1}

NMR (CDCl_3 , δ) : 2.4 (3H, s), 5.25 (2H, s), 6.77-8.05 (11H, m)

Analysis: Calcd. for $\text{C}_{18}\text{H}_{16}\text{N}_4$

Calcd.: C 74.98 H 5.59 N 19.43

Found : C 74.73 H 5.60 N 19.41

Example 10

2-(Imidazol-1-yl)methyl-1-(4-methoxyphenyl)-1H-benzimidazole was obtained according to substantially the same manner as that of Example 1.

mp. 113-114°C

IR (Nujol) : 1612, 1585, 1512 cm^{-1}

NMR (CDCl_3 , δ) : 3.9 (3H, s), 5.25 (2H, s), 6.8-8.0 (11H, m)

Analysis: Calcd. for $C_{18}H_{16}N_4O$

Calcd.: C 71.04 H 5.30 N 18.41

Found : C 71.06 H 5.38 N 18.39

5 Example 11

1-(4-Chlorophenyl)-2-(imidazol-1-yl)methyl-1H-benzimidazole was obtained according to substantially the same manner as that of Example 1.

mp. 121-122°C

10 IR (Nujol) : 1607, 1490 cm^{-1}

NMR ($CDCl_3$, δ) : 5.25 (2H, s), 6.75-8.0 (11H, m),

Analysis: Calcd. for $C_{17}H_{13}ClN_4$

Calcd.: C 66.13 H 4.24 N 18.15

Found : C 66.42 H 4.33 N 18.17

15

Example 12

2-(Imidazol-1-yl)methyl-5-methoxy-1-phenyl-1H-benzimidazole was obtained according to substantially the same manner as that of Example 1.

20 mp. 104-106°C

IR (Nujol) : 1620, 1597, 1500 cm^{-1}

NMR ($CDCl_3$, δ) : 3.85 (3H, s), 5.23 (2H, s),

6.7-7.7 (11H, m)

Analysis: Calcd. for $C_{18}H_{16}N_4O$

25 Calcd.: C 71.04 H 5.30 N 18.41

Found : C 71.00 H 5.43 N 18.30

Example 13

30 2-(Imidazol-1-yl)methyl-1-methyl-1H-benzimidazole was obtained according to substantially the same manner as that of Example 1.

mp. 151-152°C

IR (Nujol) : 1618, 1590, 1510 cm^{-1}

NMR ($CDCl_3$, δ) : 3.57 (3H, s), 5.35 (2H, s),

35 6.85-7.95 (7H, m)

Analysis: Calcd. for $C_{12}H_{12}N_4$

Calcd.: C 67.91 H 5.70 N 26.40

Found : C 68.34 H 5.80 N 26.42

5 Example 14

A mixture of 2-chloromethyl-1-phenyl-1H-benzimidazole (4.8 g) and 1-acetyl-4-methylimidazole (2.4 g) in acetonitrile (25 ml) was refluxed for 48 hours. After removal of the solvent, the residue was pulverized in diethyl ether to give powder, which was taken up in 2.5N sodium hydroxide (20 ml) and refluxed for 15 minutes. The resultant mixture was extracted twice with chloroform. The combined organic layer was extracted twice with 1N hydrochloric acid. The combined aqueous layer was adjusted to pH 8.0 with aqueous sodium bicarbonate and extracted twice with chloroform. The combined extract was washed with water, dried over magnesium sulfate and evaporated under reduced pressure. The resultant residue was subjected to column chromatography on silica gel (60 g) eluting with a mixture of chloroform and methanol (50:1) to give an oily product. The oily product was triturated twice in diethyl ether and then in the mixture of diethyl ether and isopropyl alcohol (2:1) to give powder of 2-(5-methylimidazol-1-yl)methyl-1-phenyl-1H-benzimidazole (0.65 g).

mp. 100-102°C

IR (Nujol) : 1610, 1595, 1570, 1500 cm^{-1}

NMR ($CDCl_3$, δ) : 1.95 (3H, s), 5.15 (2H, s),
6.65 (1H, brs), 6.8-8.0 (10H, m)

30 Analysis: Calcd. for $C_{18}H_{16}N_4$

Calcd.: C 74.98 H 5.59 N 19.43

Found : C 74.86 H 5.67 N 19.28

Example 15

35 Heating of 2-(imidazol-1-yl)-N-(2-anilinopyridin-

3-yl)acetamide (150 mg) at 200°C for 5 minutes afforded a crude cyclized product, which was recrystallized from a mixture of ethyl acetate and n-hexane to give 2-(imidazol-1-yl)methyl-3-phenyl-3H-imidazo[4,5-b]pyridine (116 mg).

IR (Nujol) : 1595, 1500 cm^{-1}

NMR (CDCl_3 , δ) : 5.3 (2H, s), 6.76-7.8 (9H, m), 8.15 (1H, dd, $J=8\text{Hz}$, 2Hz), 8.4 (1H, dd, $J=6\text{Hz}$, 2Hz)

Example 16

To a solution of 2-anilinoaniline (1.84 g) and 3-pyridineacetic acid (1.37 g) in chloroform (20 ml) was dropwise added phosphorus oxychloride (1.53 g) and the mixture was refluxed for 6 hours. After removal of the solvent under reduced pressure, the resultant residue was poured into a saturated aqueous solution of sodium bicarbonate. The mixture was extracted twice with ethyl acetate. The combined extract was washed with water and a saturated aqueous solution of sodium chloride successively, and dried over magnesium sulfate. The solvent was removed under reduced pressure to give a crystalline product, which was recrystallized from a mixture of isopropyl alcohol and n-hexane (1:2) to give 2-(pyridin-3-yl)methyl-1-phenyl-1H-benzimidazole (1.0 g). mp. 115-116°C

IR (Nujol) : 1615, 1595, 1575, 1500 cm^{-1}

NMR (CDCl_3 , δ) : 4.2 (2H, s), 6.9-8.0 (11H, m), 8.25 (1H, d, $J=2\text{Hz}$), 8.45 (1H, dd, $J=6\text{Hz}$, 2Hz),

Analysis: Calcd. for $\text{C}_{19}\text{H}_{15}\text{N}_3$

Calcd.: C 79.98 H 5.30 N 14.73

Found : C 79.96 H 5.52 N 14.68

Example 17

2-(Pyridin-2-yl)methyl-1-phenyl-1H-benzimidazole

was obtained according to substantially the same manner as that of Example 16.

mp. 106-108°C

IR (Nujol) : 1590, 1570, 1500 cm^{-1}

5 NMR (CDCl_3 , δ) : 4.4 (2H, s), 6.9-8.0 (12H, m),
8.3-8.6 (1H, m)

Mass: 285 (M^+)

Example 18

10 3-(1-Phenyl-1H-benzimidazol-2-yl)methyl-
benzothiazolin-2-one was obtained according to substantially the same manner as that of Example 16.

mp. 196-198°C

IR (Nujol) : 1678, 1595, 1500 cm^{-1}

15 NMR (CDCl_3 , δ) : 5.4 (2H, s), 6.8-7.9 (13H, m),

Analysis: Calcd. for $\text{C}_{21}\text{H}_{15}\text{N}_3\text{OS}$

Calcd.: C 70.57 H 4.23 N 11.76

Found : C 70.62 H 4.38 N 11.77

20 Example 19

2-[2-(Imidazol-1-yl)ethyl]-1-phenyl-1H-
benzimidazole monohydrate was obtained according to substantially the same manner as that of Example 16.

mp. 80-84°C

25 IR (Nujol) : 1610, 1595, 1500 cm^{-1}

NMR (CDCl_3 , δ) : 3.2 (2H, t, $J=6\text{Hz}$), 4.5 (2H, t,
 $J=6\text{Hz}$), 6.7-8.0 (12H, m)

Analysis: Calcd. for $\text{C}_{18}\text{H}_{16}\text{N}_4\cdot\text{H}_2\text{O}$

Calcd.: C 70.57 H 5.92 N 18.29

30 Found : C 71.03 H 6.03 N 18.38

Example 20

To a solution of 3-amino-2-anilinopyridine (2.0 g)
and 2-(imidazol-1-yl)acetic acid (1.56 g) in pyridine
35 (20 ml) was dropwise added phosphorus oxychloride (1.91 g)

and the mixture was refluxed for 30 minutes. The reaction mixture was poured into saturated aqueous sodium bicarbonate and extracted three times with ethyl acetate. The combined extract was washed with saturated aqueous sodium chloride and dried over magnesium sulfate. The solvent was removed under reduced pressure to give a residue, which was subjected to column chromatography on silica gel (20 g) eluting with a mixture of dichloromethane and ethanol. The fractions containing the desired compound were combined and evaporated in vacuo. The resultant crystal was recrystallized from a mixture of ethyl acetate and n-hexane to give 2-(imidazol-1-yl)methyl-3-phenyl-3H-imidazo[4,5-b]pyridine (0.8 g).

mp. 146-148°C
IR (Nujol) : 1595, 1500, 1430 cm^{-1}
NMR (CDCl_3 , δ) : 5.3 (2H, s), 6.76-7.8 (9H, m),
8.15 (1H, dd, J=8Hz, 2Hz), 8.4 (1H, dd,
J=6Hz, 2Hz)
Analysis: Calcd. for $\text{C}_{16}\text{H}_{13}\text{N}_5$
Calcd.: C 69.80 H 4.76 N 25.44
Found : C 70.24 H 5.11 N 25.63

Example 21

A mixture of 2-cyanomethyl-1-phenyl-1H-benzimidazole (6.2 g), sodium azide (5.18 g) and ammonium chloride (4.27 g) in N,N-dimethylformamide (62 ml) was heated at 100°C for 4 hours. The reaction mixture was poured into water (620 ml) and the aqueous mixture was adjusted to pH 9 with 1N sodium hydroxide. After being washed with ethyl acetate, the aqueous layer was further adjusted to pH 4 with acetic acid and extracted twice with ethyl acetate. The combined extract was washed with water and saturated sodium chloride in turn and dried over magnesium sulfate. The solvent was removed under reduced

pressure to give a residue, which was recrystallized from a mixture of methanol and diisopropyl ether to give 5-(1-phenyl-1H-benzimidazol-2-yl)methyl-1H-tetrazole (5.3 g).

5 mp. 185-187°C (dec.)
IR (Nujol) : 1595, 1570, 1500, 1465 cm^{-1}
NMR (DMSO- d_6 , δ) : 4.6 (2H, s), 7.1-7.9 (9H, m)
Analysis: Calcd. for $\text{C}_{15}\text{H}_{12}\text{N}_6$
10 Calcd.: C 65.21 H 4.38 N 30.42
Found : C 65.06 H 4.58 N 30.24

Example 22

A mixture of 2-cyanomethyl-1-phenyl-1H-benzimidazole (1.8 g), dicyanodiamide (785 mg) and potassium hydroxide (87 mg) in methyl cellosolve was refluxed for 2 hours. After the mixture was cooled, the resultant precipitates were filtered off and washed with water and methanol in turn to give 2,4-diamino-6-(1-phenyl-1H-benzimidazol-2-yl)methyl-1,3,5-triazine (1.8 g).

20 mp. 337-340°C
IR (Nujol) : 3300, 3100, 1665, 1635 cm^{-1}
NMR (CF_3COOH , δ) : 4.8 (2H, brs), 7.2-8.2 (9H, m),
25 Analysis: Calcd. for $\text{C}_{17}\text{H}_{15}\text{N}_7$
Calcd.: C 64.34 H 4.76 N 30.90
Found : C 64.28 H 4.88 N 31.02

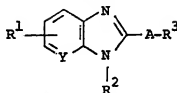
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-1-

Patent Claims

1. A compound of the formula:

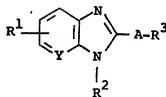


wherein A is a lower alkylene radical, R^1 is a hydrogen
 5 or halogen atom or a lower alkyl or lower alkoxy
 radical, R^2 is a hydrogen atom or a lower alkyl,
 cyclo(lower)alkyl, pyridyl, ar(lower)alkyl, which may
 be substituted by halogen, or aryl radical, which may
 be substituted by lower alkyl, lower alkoxy, hydroxy
 10 or halogen, R^3 is a nitrogen-containing unsaturated
 heterocyclic radical, which may be substituted by
 lower alkyl or amino, and Y is =C- or =N-; and the
 pharmaceutically-acceptable salts thereof.

2. A compound according to claim 1, wherein R^1 is
 15 a hydrogen atom or a lower alkoxy radical and R^2 is
 a hydrogen atom or a lower alkyl, ar(lower)alkyl,
 which may be substituted by halogen, or aryl radical,
 which may be substituted by lower alkyl, lower alkoxy
 or halogen.

20 3. A compound according to claim 2, wherein R^3 is
 an imidazolyl, pyridyl, triazolyl, tetrazolyl, 1,3,5-
 triazinyl or 2-oxo-benzothiazoliny radical, which
 may be substituted by lower alkyl or amino, and Y is
 =C-.

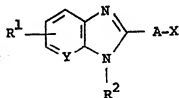
4. A compound according to claim 3, wherein R^1 is a hydrogen atom, R^2 is an ar(lower)alkyl radical, which may be substituted by halogen, or an aryl radical, which may be substituted by lower alkoxy or halogen, and R^3 is a pyridyl or imidazolyl radical, optionally substituted by lower alkyl.
5. A compound according to claim 4, wherein A is a methylene radical, R^2 is a phenyl radical and R^3 is an imidazolyl radical, optionally substituted by lower alkyl.
6. A compound according to claim 2, wherein R^3 is an imidazolyl radical and Y is =N-.
7. 2-(Imidazol-1-yl)-methyl-1-phenyl-1H-benzimidazole.
8. A process for preparing a compound of the formula:



- wherein A is a lower alkylene radical, R^1 is a hydrogen or halogen atom or a lower alkyl or lower alkoxy radical, R^2 is a hydrogen atom or a lower alkyl, cyclo(lower)alkyl, pyridyl, ar(lower)alkyl, which may be substituted by halogen, or aryl radical, which may be substituted by lower alkyl, lower alkoxy, hydroxy or halogen, R^3 is an N-containing unsaturated heterocyclic radical, which may be substituted by lower alkyl or amino, and Y is =C- or =N-; and the

pharmaceutically acceptable salts thereof, which comprises

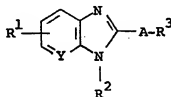
a) reacting a compound of the formula:



- 5 wherein A, R¹, R² and Y have the same meanings as above and X is an acid residue or a salt thereof with a compound of the formula:

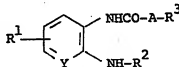


- wherein R³ has the same meaning as above, or with a
10 salt thereof to give a compound of the formula:

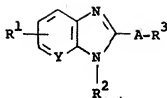


or a pharmaceutically-acceptable salt thereof, wherein R¹, R², R³, A and Y have the same meanings as above; or
b) cyclising a compound of the formula:

15

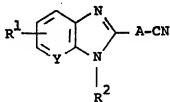


wherein A, R¹, R², R³ and Y have the same meanings as above, or a salt thereof to give a compound of the formula:

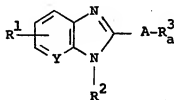


or a pharmaceutically-acceptable salt thereof,
wherein R^1 , R^2 , R^3 , A and Y have the same meanings
as above; or

- 5 c) reacting a compound of the formula:



wherein A, R^1 , R^2 and Y have the same meanings as
above, or a salt thereof with dicyanodiamide or a
salt thereof or an azide compound to give a compound
10 of the formula:



or a pharmaceutically-acceptable salt, wherein R^1 ,
 R^2 , A and Y have the same meanings as above and R_a^3
is a 1H-tetrazol-5-yl or 2,4-diamino-1,3,5-triazin-
15 6-yl radical.

9. A pharmaceutical composition comprising a compound of claim 1, as an effective ingredient, in association with a pharmaceutically-acceptable, substantially non-toxic carrier or excipient.
- 5 10. A method for treating ulcers, which comprises administering an effective amount of a compound of claim 1 to a human or to an animal.

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